

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 0 460 426 B1

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention
of the grant of the patent:
17.09.1997 Bulletin 1997/38

(51) Int Cl.⁶: C07K 1/19, C07K 16/00,
C12P 21/00

(21) Application number: 91107661.0

(22) Date of filing: 11.05.1991

(54) Protein purification method
Proteinreinigungsmethode
Méthode de purification des protéines

(84) Designated Contracting States:
AT BE CH DE DK ES FR GB GR IT LI LU NL SE

• Lee, Vivian W.
Richmond, California (US)

(30) Priority: 25.05.1990 US 528523

(74) Representative: Kirchner, Dietrich, Dr. et al
Bayer AG,
Konzernbereich RP
Patente und Lizenzen
51368 Leverkusen (DE)

(43) Date of publication of application:
11.12.1991 Bulletin 1991/50

(73) Proprietor: Bayer Corporation
Pittsburgh, PA 15219-2502 (US)

(56) References cited:
EP-A- 0 209 786 EP-A- 0 239 565
WO-A-89/03840

(72) Inventors:
• Coan, Michael H.
El Cerrito, California (US)

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent granted. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

BEST AVAILABLE COPY

EP 0 460 426 B1

Description

Background of the Invention

Field:

This disclosure is concerned generally with a method of purifying an antibody and specifically with a three step method involving pH adjustments to purify the antibody without the use of precipitation agents or salt solutions.

Prior Art:

Current methods for purification of proteins include using precipitation agents such as salts or non-salt substances such as polyethylene glycol (PEG). Unfortunately, in many cases the protein yields are less than desired. The methods are often time consuming and often require the use of specialized equipment such as large centrifuges. Also, it is often difficult to scale up a precipitation method and even if this can be done dissolution of the resulting precipitate can be slow and is not always complete.

Another common method for purifying proteins includes passing a protein-containing solution over or through an appropriate ion exchange material at a solution pH which facilitates binding of the desired protein. This is commonly followed by a washing step and an elution step at a different ionic strength or pH which facilitates the release of the protein. See, for example, Lamy, J. et al, Arch Biochem. Biophys. 193 pp 140-149 (1979), showing that protein can be eluted from DEAE Sepharose® by using a descending pH gradient. In general, altering the pH of a protein towards its isoelectric point causes it to lose net charge and elute from an ion exchanger.

Elution of materials from ion exchangers by pH change and ionic strength change is described more fully in Morris et al, Separation Methods in Biochemistry Pittman and Sons, London, 1964. See also Robert Scopes, Protein Purification, Springer-Verlag, New York, NY, 1982. At page 85, the author points out that the use of a change in buffer pH is generally not very successful (for protein purification).

EP-A-0 209 786 relates to a process of separating hTNF from a mixture of N-terminal methionine containing hTNF using a cation exchanges containing sulphonyl- or sulphaalkyl groups.

Pharmacia Ion Exchange Chromatography booklet 1983 (pages 45 - 47) represents state of the art protein binding and elution techniques and describes a two-step process for eluting a bound protein (arthropod haemocyanin) from an ion exchanger (DEAE-Sepharose (L-6B)).

Cellulosic ion exchangers, Ed. E.A. Peterson, 1980, pages 262 - 264 describes a general concept of the purification of proteins with ion-exchange chromatogra-

phy.

The reference Handbook "Sorbent Extraction Technology" Analytichem International, Ed. K.C. van Horne, 1985, pages 29 - 32 and 59 - 60 is disclosing in Section 4.3 the relationship between the charged isolate molecule (which means the molecule to be bound) and the sorbent carrying a charge opposite to that of the isolate (e.g. an ion exchanger). Section 10.2.2. of said reference refers to wash solvents which do not elute the isolate.

We have now found that by slightly modifying the above ion exchange/pH methods, we can purify an antibody such as a monoclonal antibody in a relatively simple manner that avoids protein loss commonly associated with existing protein purification methods, especially methods based on the use of solutions of increasing ionic strength. Details of our method are described below.

SUMMARY OF THE INVENTION

Our method of purifying an antibody from an aqueous solution containing impurities comprises three essential steps. In the first step, the solution is contacted with an ion exchange material at a solution pH which facilitates binding of the desired antibody (first pH). Other substances (impurities) may also be bound during this step. In the next step the exchange material to which the antibody is bound is washed with a solution having a different pH which does not elute the protein (second pH). This pH is one which does not facilitate binding of the unbound protein to the ion exchange material but, for some reason, does not facilitate elution of the already bound protein. In the third step, the antibody is eluted with a solution having a pH which facilitates protein elution (third pH).

In the preferred embodiment, the antibody is an antibody which binds to the tumor necrosis factor. In the examples below, the method of this disclosure is illustrated using a monoclonal antibody that specifically binds to a substance known as tumor necrosis factor or TNF. In those examples we found that best results could be obtained by binding, washing and then eluting the antibody at three distinct and increasing values of pH.

SPECIFIC EMBODIMENTS

Our disclosure is further illustrated in the examples below in which we purified anti-TNF antibodies expressed from a sub-clone of a deposited cell line having an ATCC Accession No. HB9736. This cell line is described further in Patent Application S.N. 220,206, filed July 18, 1988, incorporated herein.

The improved purification method of this disclosure was discovered while trying to find a way to reduce protein loss that accompanied the use of a precipitation step to purify the monoclonal antibodies to TNF in a clarified (filtered) tissue culture fluid (TCF).

BEST AVAILABLE COPY

In the TCFs we worked with, the major protein components were the anti-TNF monoclonal antibodies and human albumin, the albumin being present to stabilize the cells of the original (non-clarified) TCF. It had been found particularly difficult to separate albumin from the anti-TNF because of the similarity of their isoelectric points (only 1.5 pH unit difference). Other antibodies are generally easier to separate from albumin because they have a higher isoelectric point (more than 2.0 pH units). As used herein, a significant difference in isoelectric point between human albumin and an antibody, especially a monoclonal antibody, refers to a difference of more than about 2.0 pH units. Having similar isoelectric points means that the difference is less than 2.0 pH units.

We decided to try to replace an unsatisfactory PEG precipitation step with a cation exchange process even though that type of process is not more commonly used for protein purification.

The initial process we used was a fairly standard ion exchange method. That is, at a particular pH and ionic strength we equilibrated a cation exchange column. The ion exchange matrix was S-Sepharose® which is sulphopropyl surface modified agarose supplied by Pharmacia. The aqueous feed solution (clarified TCF) containing the desired protein was adjusted to the same conditions (pH 4.6, 0.01 M sodium acetate) and applied to the column. TCF is the harvest fluid from the cell fermenter. It contains carbohydrates, salts, amino acids, proteins, other cell growth factors, and cells. Clarified TCF has had the cells removed by filtration or some other means. The clarified TCF may be concentrated. We determined the capacity of the column at these conditions and then loaded it to capacity or below. Then, after washing the column with equilibration buffer, the column was eluted, as is typical in ion exchange, by increasing ionic strength. In this case, the conditions were pH 4.6, 0.01 M sodium acetate, 0.27 NaCl (i.e. the salt concentration was raised).

Using that technique, the recovery of antibody was >85% with about 15% purity. The next step in this process would have been to use an anion exchange material (e.g., Q-Sepharose®) for further purification.

It then occurred to us that the elution of the S-Sepharose® column with a high salt concentration created the necessity of adding a new step (salt reduction or removal) where more protein loss would be expected to occur. This would be undesirable.

Instead of following the above elution method (changing ionic strength), we decided to consider conditions of washing and elution which did not increase the ionic strength but, instead, increased the pH.

In our particular examples, we surprisingly found a set of conditions which would enable much of the impurities (mostly albumin in our examples) to be washed off. In another step, by raising the pH again, we were able to elute the antibody. Thus, in our examples, we were able to find that the anti-TNF should be loaded at an

initial value, then washed at a different pH value (in this case higher) and then eluted at yet a third pH value which in this case was the highest.

Surprisingly, we found that if the column was equilibrated and loaded at the intermediate pH, little or no binding of the protein occurred. Although the exact mechanism is not clear, it is thought that conformational changes in the protein at different pH conditions may affect binding. In the case of our examples, it is thought that the lowest pH brings about a conformational or conformational change of the anti-TNF which facilitates binding to the ion exchange matrix. At the intermediate pH, the unbound protein may be in a conformation in which the binding sites are blocked due to a conformational change caused by that pH. If, however, the anti-TNF is already bound to the matrix, the conformational change caused by the intermediate pH may not be possible.

As used herein, the expression different pH, when applied to the intermediate washing step of this disclosure and the claims below, refers to a pH at which a protein (such as the anti-TNF monoclonal antibodies of the examples below) will not bind to an ion exchange material (such as S-Sepharose®) if in the free or unbound state but which will not elute from the same ion exchange material if already bound to it.

Example 1

An S-Sepharose® (Pharmacia) column was equilibrated with 0.01 M sodium acetate at pH 4.6. The anti-TNF containing solution (TCF) was then applied to the column as next described. The solution was diluted with water (approximately 1 part water to 2 parts TCF). The pH of the diluted TCF was adjusted to pH 4.6 with 1M acetic acid. The absorbance at 280 nm (A_{280}) was measured and 20 A_{280} units were applied per mL of column. In these examples, we define a protein concentration of one unit per mL as having an A_{280} of 1. The column was washed with the equilibration buffer until the eluting absorbance returned to baseline. The column was then washed with buffer of 0.01 sodium acetate, 0.05 M sodium chloride, pH 5.5. Some proteins eluted, and the column was washed until the A_{280} returned to baseline. In the final step, the antibodies were eluted at a solution pH of 6.5. The antibody recovery was 90% with a purity of 30%.

Example 2

The same starting clarified TCF as in Example 1 was used. This time, however, the S-Sepharose® column was equilibrated with 0.01 sodium acetate, 0.05 M NaCl, pH 5.5. Also, the TCF was diluted as in Example 1, but was then adjusted to pH 5.5 with 1M acetic acid. The column loading was the same as described in Example 1. The column was washed until the absorbance at 280 nm returned to baseline. The elution was as in

Example 1. The recovery of antibody was only 10%. The remaining antibody was accounted for in the unbound fractions.

Given the above disclosure, it is thought that variations will occur to those skilled in the art of protein purification. Accordingly, it is intended that the above examples should be construed as illustrative only and that the scope of the invention of this disclosure should be limited only to the following claims.

Claims

1. A method of purifying an antibody in an aqueous solution comprising the three essential steps of

(a) binding the antibody to an ion exchange material at a first pH for the aqueous solution which facilitates protein binding;

(b) washing the bound antibody at a second different pH which does not facilitate binding of the unbound protein to the ion exchange material but does not facilitate elution of the already bound protein;

(c) eluting the antibody at a third pH for the elution solution which facilitates elution.

2. The method of claim 1 wherein and all steps are done without substantial changes in ionic strength among the solutions.

3. The method of claim 2 wherein the antibody binds tumor necrosis factor.

4. The method of claim 1 wherein the steps use a sequence of increasing pH in going from the initial to the final step.

5. The method of claim 3 wherein the antibody is bound to the ion exchange material at a pH of about 4.6, the washing step is at a pH of about 5.5; and the elution step is at a pH of about 6.5.

6. The method of claim 5 wherein the antibody is expressed from the cell line HB 9736.

Patentansprüche

1. Verfahren zur Reinigung eines Antikörpers in einer wässrigen Lösung, umfassend die drei wesentlichen Stufen, in denen man:

(a) den Antikörper an ein Ionenaustauschmaterial bei einem ersten pH-Wert der wässrigen Lösung bindet, welcher die Proteinbindung er-

leichtert;

(b) den gebundenen Antikörper bei einem zweiten anderen pH-Wert wäscht, welcher die Bindung von ungebundenem Protein an das Ionenaustauschmaterial nicht erleichtert, die Elution des bereits gebundenen Protein aber auch nicht erleichtert;

(c) den Antikörper bei einem dritten pH-Wert der Elutionslösung eluiert, welcher die Elution erleichtert.

2. Verfahren gemäß Anspruch 1, wobei alle Stufen ohne wesentliche Änderungen bei der Ionenstärke der Lösungen durchgeführt werden.

3. Verfahren gemäß Anspruch 2, worin der Antikörper Tumornekrosefaktor bindet.

4. Verfahren gemäß Anspruch 1, wobei in den Stufen eine Abfolge steigender pH-Werte beim Gang von der Anfangs- zur Endstufe angewandt wird.

5. Verfahren gemäß Anspruch 3, wobei der Antikörper an das Ionenaustauschmaterial bei einem pH von ca. 4,6 gebunden, die Waschstufe bei einem pH von ca. 5,5 und die Elutionsstufe bei einem pH von ca. 6,5 durchgeführt werden.

6. Verfahren gemäß Anspruch 5, worin der Antikörper aus der Zelllinie HB 9736 exprimiert ist.

Revendications

1. Procédé de purification d'un anticorps dans une solution aqueuse comprenant les trois étapes essentielles suivantes :

(a) la liaison de l'anticorps à une substance échangeuse d'ions à un premier pH pour la solution aqueuse qui facilite la liaison de la protéine;

(b) le lavage de l'anticorps lié à un second pH différent qui ne facilite pas la liaison de la protéine non liée à la substance échangeuse d'ions mais ne facilite pas l'élution de la protéine déjà liée;

(c) l'élution de l'anticorps à un troisième pH pour la solution d'élution qui facilite l'élution.

2. Procédé selon la revendication 1, dans lequel toutes les étapes sont effectuées sans changement substantiel dans la force ionique des solutions.

3. Procédé selon la revendication 2, dans lequel l'anticorps se lie au facteur de nécrose tumorale.

BEST AVAILABLE COPY

4. Procédé selon la revendication 1, dans lequel les étapes utilisent une séquence de pH croissants en allant de l'étape initiale à l'étape finale.
5. Procédé selon la revendication 3, dans lequel l'anticorps est lié à la substance échangeuse d'ions à pH d'environ 4,6, l'étape de lavage est à pH d'environ 5,5; et l'étape d'élution est à pH d'environ 6,5.
6. Procédé selon la revendication 5, dans lequel l'anticorps est exprimé à partir de la lignée cellulaire HB 9736.

15

20

25

30

35

40

45

50

55

5

BEST AVAILABLE COPY